Appln. No. 09/269,321 Amendment dated March 15, 2005 Reply to Office Communication dated March 8, 2005

## **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

- 1-14. (Canceled)
- 15. (Previously presented) The method of claim 25, wherein the nucleic acid cassette is present in a viral vector or nucleic acid delivery system.
- 16. (Previously presented) The method of claim 25 wherein the malignant cell is a solid tumor.
- 17. (Previously presented) The method of claim 16 wherein the solid tumor is a glioma.
- 18. (Currently amended) The method of claim 17, wherein the nucleic acid <u>cassette</u> eassettes is present in a vector, wherein the vector is an adenovirus vector or a herpes virus vector.
- 19. (Original) The method of claim 16, wherein the nucleic acid sequence of interest encodes a negative potentiator.
- 20. (Previously presented) The method of claim 19, wherein the gene of interest is a suicide gene, a dominant negative mutant or a cytotoxin.
- 21. (Previously presented) The method of claim 20, wherein the gene of interest is a suicide gene.

BOS1473728.1 2

Appln. No. 09/269,321 Amendment dated March 15, 2005 Reply to Office Communication dated March 8, 2005

- 22. (Original) The method of claim 21, wherein the suicide gene is HSV thymidine kinase.
- 23. (Previously presented) The method of claim 20, wherein the gene of interest is a cytotoxin.
- 24. (Previously presented) The method of claim 23, wherein the cytotoxin contains at least Domain III of *Pseudomonas extoxin A*.
- 25. (Currently amended) A method of selectively expressing a gene in a malignant cell comprising:
- (a) determining whether the malignant cell expresses sufficient E2F to cause increased expression of a gene operably linked to an E2F responsive promoter when compared to a mitotically active non-malignant cell;
- (b) selectively expressing the gene in said malignant cell that was determined to express sufficient E2F by adding an effective amount of a nucleic acid cassette to the malignant cell that was determined to express sufficient E2F, wherein said nucleic acid cassette comprises an E2F responsive promoter operably linked to a gene of interest, wherein said gene encodes a protein that stimulates production or expression of a cellular product that is product, a positive potentiator or encodes a gene that inhibits production or expression of a cellular product that is product that is product, a negative potentiator; and
  - (c) waiting until the nucleic acid cassette transduces the malignant cell, eell; and

BOS1473728.1 3

Appln. No. 09/269,321 Amendment dated March 15, 2005 Reply to Office Communication dated March 8, 2005

(d) selectively expressing the gene by the E2F in said malignant cell causing the E2F responsive promoter to express said gene.

- 26. (Previously presented) The method of claim 25, wherein the E2F responsive promoter is selected from the group of promoters consisting of E2F1 promoter, dihydrofolate reductase promoter, DNA polymerase α promoter, c-myc promoter and β-myb promoter.
- 27. (Previously presented) The method of claim 25, wherein the gene of interest is selected from the group consisting of cytokines or costimulatory molecules.

28-40. (Canceled).

BOS1473728.1 4